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IN THE UNITED STATES PATENT & TRADEMARK OFFICE

RE APPLICATION OF:

VILLIAM M. CANFIELD

: EXAMINER: PATTERSON, JR., C.

SERIAL NO: 09/895,072

FILED: July 02, 2001

: GROUP ART UNIT: 1652

FOR: METHODS FOR PRODUCING

HIGHLY PHOSPHORYLATED LYSOSOMAL HYDROLASES

DECLARATION UNDER 37 C.F.R. 1.132

ASSISTANT COMMISSIONER FOR PATENTS WASHINGTON, D.C. 20231

SIR:

Now comes Stuart Kornfeld, M.D. who states that:

- I am the David C. and Betty Farrell Distinguished Professor of Medicine at
 Washington University, St. Louis, Missouri
- 2. My area of expertise is in lysosomal enzyme trafficking. My curriculum vitae is attached as Exhibit 1.
- 3. I am formerly a member of the board of directors of Novazyme Pharmaceuticals, Inc., the prior assignee of the application.

- 4. It is my understanding that a point of contention in this application is whether it would require undue experimentation to purify N-acetylglucosamine-1-phosphotransferase to a specific activity of at least 10⁷ pmol/h/mg and/or phosphodiester _-GlcNAcase to a specific activity 472,000 units/mg based on the descriptions in Bao et al ((1996) J. Biol. Chem. 271(49):31437-31445); Bao et al ((1996) J. Biol. Chem. 271(49):31446-31461) and Kornfeld ((1998) J. Biol. Chem. 273(36):23203-23210) notwithstanding the fact that the antibodies used in those publications were not made available to the public.
- 5. During a period of approximately 15 years, my lab and other labs attempted to purify these two enzymes without success notwithstanding employing state of the art biochemical methods. In addition, I am aware of other attempts at purification using monoclonal antibody affinity techniques, which also failed to yield anything other than a partially purified preparation with relatively low specific activity.
- 6. To be useful for affinity purification, the monoclonal antibodies must have a collection of specific attributes. These antibodies must bind with high affinity as the enzymes are a trace component of a crude protein preparation; the antibodies must not inhibit the intrinsic enzymatic activity while bound; and the binding between the antibody and the protein must be reversible under mild conditions consistent with the stability profile of the target enzyme. Although it is presumably possible to isolate other monoclonal antibodies with these required properties success would be an extremely rare event and as a result would require undue experimentation. Therefore,

it would require undue experimentation to purify the N-acetylglucosamine-1-phosphotransferase and/or phosphodiester _-GlcNAcase without the two specific antibodies described in the above-identified patent application.

- 7. Therefore, the descriptions in the publications of paragraph 4 above, which are also cited by the patent office, do not provide sufficient information to enable one of skill in the art to purify the phosphotransferase and the N-acetylglucosamine-1-phosphodiester _-N-Acetylglucosaminidase enzymes to the specific activities noted above.
- I declare under penalty of perjury that the foregoing is believed to be true and accurate.

Sturt Kanfold M.O.

12/19/03

Stuart Kornfeld, M.D.

Date

STUART KORNFELD

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August 28, 2002

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	Research Assistant, Biochemistry Department, Washington University School of Medicine, St. Louis, MO
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1963 - 1965	Research Associate, National Institute of Arthritis and Metabolic Diseases, NIH
1965 - 1966	Assistant Resident in Ward Medicine, Barnes Hospital, St. Louis, MO
1966 - 1967	Instructor in Medicine, Washington University School of Medicine, St. Louis, MO

- 1967 1970 Assistant Professor of Medicine, Washington University School of Medicine
- 1968 1976 Assistant Professor of Biochemistry, Washington University School of Medicine
- 1970 1972 Associate Professor of Medicine, Washington University School of Medicine
- 1972-Present Professor of Medicine, Washington University School of Medicine
 - 1973 1976 Director, Division of Oncology, Washington University School of Medicine
 - 1976 1992 Co-Director, Division of Hematology-Oncology, Washington University School of Medicine
 - 1993-Present Co-Director, Division of Hematology Washington University School of Medicine

2000-Present David C. and Betty Farrell Distinguished Professor of Medicine

Hospital Appointments:

Barnes Hospital Jewish Hospital

Appointments and Committees:

1991-1997 Director, Medical Scientist Training Program 2000-presentCo-Director, Physician Scientist Training Program

Medical Licensure and Board Certification:

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Military Service:

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Honors and Awards:

Biochemistry Award, Washington University School of Medicine, 1959 Bordon Award for Outstanding Undergraduate Research, 1962 American Cancer Society Faculty Research Associate, 1966-1971 Research Career Development Award, National Institutes of Health, 1971-

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Institute of Medicine- 1983

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Alumni/Faculty Award, Washington University School of Medicine, 1987 Jubilee Lecturer and Harden Medallist, The Biochemical Society, 1989 Passano Award, 1991 (with William Sly)

E. Donnall Thomas Lectureship and Prize, 1992

Karl Meyer Award, Society for Glycobiology, 1999

UCSD/Nature Medicine "Mentorship Award", 2002

Gerty & Carl Cori Faculty Recognition Award, Washington University,

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Editorial Responsibilities:

1972-1996	Editorial Board, Archives of Biochemistry and Biophysics		
1976-1981	Editorial Board, Journal of Biological Chemistry		
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1981-1982	Editor, Journal of Clinical Investigation		
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Professional Societies and Organizations:

American Society for Clinical Investigation American Society of Biological Chemists American Society of Hematology Association of American Physicians Foreign Member, Finnish Society of Sciences and Letters

Board Memberships:

1972-1975 Councillor, American Society for Clinical Investigation

1974-1977	Member, NIH Cell Biology Study Section
1983-1987	Member, NIADDK Board of Scientific Counselors
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